

Interplay Between Hydrogen Sulfide and Adrenergic and Muscarinic Receptors in the Mouse Atrium

Lifanova A., Khaertdinov N., Sitdikova G.

Kazan Federal University, 420008, Kremlevskaya 18, Kazan, Russia

Abstract

© 2016, Springer Science+Business Media New York. Hydrogen sulfide (H₂S) is synthesized endogenously, and it has a negative inotropic effect on myocardium of different animal species. Interaction between H₂S and adrenergic and muscarinic receptors in regulation of mouse atrium contractile function was investigated in this study. Sodium hydrosulfide (NaHS, 300 μM), the H₂S donor, decreased the contraction force of atrium. NaHS did not affect the positive inotropic effect of β-adrenoceptors (β-AR) activation by isoproterenol (ISO, 1 μM). The effect of the H₂S donor under β-AR stimulation showed no differences comparison to control values. The agonist of muscarinic receptors carbachol (1 μM) induced a negative inotropic effect and partially prevented the reduction of cardiac muscle contractility by NaHS. Moreover, the effect of carbachol was more pronounced after preliminary application of NaHS. At the same time, after inhibition of β-AR (propranolol, 1 μM) or muscarinic receptors (atropine, 1 μM), negative inotropic effect of NaHS was the same as in control conditions. These results suggest that the H₂S effects are mediated by intracellular signaling pathways activated by muscarinic receptors.

<http://dx.doi.org/10.1007/s12668-016-0355-1>

Keywords

Hydrogen sulfide (H₂S), Mouse atrium, Muscarinic receptors, β-Adrenoceptors

References

- [1] Sitdikova, G. F., & Zefirov, A. L. (2010). Hydrogen sulfide: from the Parisian sewer system to a signaling molecule. *Priroda (Rus.)*, 9, 29–37.
- [2] Elsey, D., Fowkes, R., Baxter, G. (2010). Regulation of cardiovascular cell function by hydrogen sulfide (HS). *Cell Biochemistry and Function*, 28(2), 95–106. doi:10.1002/cbf.1618.
- [3] Yong, Q. C., Pan, T. T., Hu, L. F., Bian, J. S. (2008). Negative regulation of beta-adrenergic function by hydrogen sulfide in the rat hearts. *Journal of Molecular and Cellular Cardiology*, 44(4), 701–710. doi:10.1016/j.yjmcc.2008.01.007.
- [4] Khaertdinov, N. N., Lifanova, A. S., Gizzatullin, A. R., Sitdikova, G. F. (2015). Role of K(ATP)-channels in the effects of hydrogen sulfide on the contractility of rat ventricular myocardium. *Genes and Cells*, 10(4), 103–105.
- [5] Lifanova, A. S., Khaertdinov, N. N., Zakharov, A. V., Gizzatullin, A. R., Sitdikova, F. G. (2014). Role of potassium channels in the negative inotropic effect of hydrogen sulfide in mouse atrium. *Genes and Cells*, 9(3), 94–98.
- [6] Sitdikova, G. F., Khaertdinov, N. N., Zefirov, A. L. (2011). Investigation of the role of calcium and potassium channels in hydrogen sulfide effects on myocardial contractility in frog. *Bulletin of Experimental Biology and Medicine (Rus.)*, 151(2), 124–128.

- [7] Khaertdinov, N. N., Ahmetshina, D. R., Zefirov, A. L., Sitdikova, G. F. (2013). Hydrogen sulfide in regulation of frog myocardium contractility. *Biochemistry (Moscow) Supplement Series A: Membrane and Cell Biology*, 7(1), 52-57. doi:10.1134/S1990747812030117.
- [8] Colette, C., Papapetropoulos, A., Erdelyi, K., Olah, G., Módis, K., Panopoulos, P., et al. (2012). Hydrogen sulfide and nitric oxide are mutually dependent in the regulation of angiogenesis and endothelium-dependent vasorelaxation. *Proceedings of the National Academy of Sciences of the United States of America*, 109(23), 9161-9166. doi:10.1073/pnas.1202916109.
- [9] Sekiguchi, F., & Kawabata, A. (2013). T-type calcium channels: functional regulation and implication in pain signaling. *Journal of Pharmacological Sciences*, 122(4), 244-250.
- [10] Harvey, R. D., & Belevych, A. E. (2003). Muscarinic regulation of cardiac ion channels. *British Journal of Pharmacology*, 139(6), 1074-1084.
- [11] Yamamoto, S., Miyamoto, A., Kawana, S., Namiki, A., Ohshika, H. (1998). Role of nitric oxide production through M2-cholinergic receptors in cultured rat ventricular myocytes. *Biochemical and Biophysical Research Communications*, 251(3), 791-795.